

Name: Jennifer Evans

CONFIRMATION?

Date: 6/19/14

081

WEEKLY ONE SENTENCE: continue testing RBS primer /
primer pairs

GEL LANE ORDER:

DAILY ONE SENTENCE: watch Lambert High School's 'gen
presentation, colony PCR at yesterday's
transformation - grow cultures if they work

GEL PICTURE / OTHER SPACE:

METHOD(S) OF CHOICE:

~~colony PCR~~ eye balls
colony PCR

EXPECTED RESULT:

band ~ ~~1 kb~~ 1 kb

(☒ check if APE file exists on Drive)

TEMPLATE(S): PH1/RM1 memory + PH2/RL1 memory

OLIGO(S): 080 - 085
0863

→ 081

THERMOCYCLER SETTINGS:

Haylee → vol PCR

CONTROLS:

POSITIVE:

NEGATIVE:

no polymerase

Name: Kathryn Fitton

Date: 6/19/14

WEEKLY ONE SENTENCER:

Continue testing promoter RBS primers

DAILY ONE SENTENCER:

do colony PCR ~~on~~ ^{w/} colonies ~~on~~ on plates,
grow bacteria cultures for colonies that work,
METHOD(S) OF CHOICE: listen to Lamberts presentation
colony PCR

gel

EXPECTED RESULT:

band @ 1kb

(☒ check if APE file exists on Drive)

TEMPLATE(S): p11 R11 mcherry + p12 R11 mcherry

OLIGO(S): 80-853363

THERMOCYCLER SETTINGS:

CONTROLS:

POSITIVE:

NEGATIVE:

No polymerase

CONFIRMATION?

gel

GEL LANE ORDER:

GEL PICTURE / OTHER SPACE:

Name:

Stefanos Tassoulas

CONFIRMATION?

Date:

06/29/14

WEEKLY ONE SENTENCER:

Make competent cells

GEL LANE ORDER:

DAILY ONE SENTENCER:

Work on funding

Prep. ~~Make~~ comp cells

Work on funding

METHOD(S) OF CHOICE:

- Incubator,

- Plaque

- Computer

EXPECTED RESULT:

- Success

- Sufficient cell growth

(☐ check if APE file exists on Drive)

TEMPLATE(S):

OLIGO(S):

THERMOCYCLER SETTINGS:

CONTROLS:

POSITIVE:

NEGATIVE:

GEL PICTURE / OTHER SPACE:

Name: Nishi Chaudhury

Date: 6/19/14

CONFIRMATION?
~~from~~ Get Trick gel.

WEEKLY ONE SENTENCER:

Continue the primer & promoter combination testing. ✓

GEL LANE ORDER:

1x6

4H113

4H2B

2M1B

2M2B

BL1B

RUN

2x1

PL1

PL1

PL1

PL1

PL1

PL1

GEL PICTURE / OTHER SPACE:

140619 promoter + RBS + melbary

DAILY ONE SENTENCER: - Critique/Watch Lambert High's iSEM ppt. ✓
- Miniprep the colonies taken & I made yesterday (RBS+melbary)
- Then if the concentrations are high we will create a frozen stock of them
METHOD(S) OF CHOICE: - Sequence the miniprep sample
- iSEMube 11

✓
✓
- set up another cell culture with left over bacterial cultures from miniprep.
- PCR remaining stuff with promoters.

EXPECTED RESULT:

- High concentrations in the nanodrop

✓
check if APE file exists on Drive)

TEMPLATE(S): PL1 + all RBS + melbary. ✓

OLIGO(S): 0363 2 to PL1 (84)

THERMOCYCLER SETTINGS: 94-2', 294'-15", 65-15", 68-1.5' 2x31 - 68°C-4'; 4°C ✓

CONTROLS:

POSITIVE: with polymerase

NEGATIVE: No polymerase

Name: Coleen Tran

Date: 6/19/14

WEEKLY ONE SENTENCER:

working on RBS+promoter combinations

DAILY ONE SENTENCER:

make fusion spots & sequence RBS+adhens

METHOD(S) OF CHOICE:

sequencing

insert promoter primers 146619 promoter+RBS+adhens

EXPECTED RESULT:

~1000 bp

(☒ check if APE file exists on Drive)

TEMPLATE(S):

OLIGO(S): 81, 363

THERMOCYCLER SETTINGS:

94-21; { 94'-15''; 65-15''; 68-1.5; { x31; 68-4''; 4°C

POSITIVE:

?

NEGATIVE:

no polymerase

CONFIRMATION?

GEL LANE ORDER:

Thick gel?

2 RM1 RM2 RM1 RM2 RM4 RM11N

RM2 RM2 RM2 RM2 RM2 RM2

GEL PICTURE / OTHER SPACE: